



Contains No CBI

8EHQ-0893-12199

2030 DOW CENTER
August 31, 1993

The Dow Chemical Company
Midland, Michigan 48674

A

CERTIFIED MAIL--RETURN RECEIPT
REQUESTED

Document Processing Center (TS-790)
Office of Toxic Substances
U.S. Environmental Protection Agency
401 M Street, SW
Washington, D.C. 20460
Attn: 8(e) Coordinator



8EHQ-93-12199
INIT 09/03/93

Re: Diethylene Triamine (DETA)



88930000421

Dear Sir/Madam:

The following information is being submitted by The Dow Chemical Company (Dow) pursuant to current guidance issued by EPA indicating EPA's interpretation of Section 8(e) of the Toxic Substance Control Act. Dow has made no determination as to whether a significant risk of injury to health or the environment is actually presented by the findings.

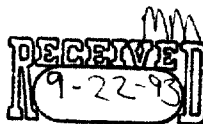
The preliminary results of a non-definitive, reproductive screen (OECD Guideline 421) have been communicated to Dow by a foreign subsidiary. The screen was conducted using Wistar rats. The animals received doses at 0, 30, 100 and 300 mg/kg/day via oral gavage. The preliminary results indicated a decrease in food consumption and decreased body weight gain in both parental males and females at the 300 mg/kg/day dose. Also, at the 300 mg/kg/day dose, the results indicated a decreased litter size, increased post-implantation loss and an increase in the length of gestation. At the 100 mg/kg/day dose, an increase in the length of gestation and a non-statistically significant increase in the post-implantation loss were observed. The statistically significant increase in the length of gestation observed at the 100 and 300 mg/kg/day doses may have been spurious as the length of gestation in these two groups of animals was within the range considered normal (21-22 days). No significant parental or reproductive/developmental effects were observed at the 30 mg/kg/day dose.

No histopathologic evaluation or report is available at this time.

Sincerely,

Paul A. Wright

Paul A. Wright
Senior Attorney
Legal Department
517/636-1853



93SEP-3 PM 1:50



UNITED STATES ENVIRONMENTAL PROTECTION AGENCY
WASHINGTON, D.C. 20460

Paul A. Wright
Attorney, Legal Department
The Dow Chemical Company
2030 Dow Center
Midland, Michigan 48674

OFFICE OF
PESTICIDES AND TOXIC
SUBSTANCES

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This letter formally acknowledges EPA's receipt of information submitted by your organization under Section 8(e), the "substantial risk" information reporting provision of the Toxic Substances Control Act (TSCA). For your reference, copies of the first page(s) of your submission(s) are enclosed and display the TSCA Section 8(e) Document Control Number (i.e., 8EHQ-0000-0000 Init.) assigned by EPA to your submission(s). Please refer to this cited number when submitting follow-up or supplemental information.

Please note that all submitted correspondence will be placed in the public files unless confidentiality is claimed according to the procedures outlined in Part X of EPA's TSCA Section 8(e) policy statement (43 FR 11110, March 16, 1978).

Confidential submissions submitted pursuant to the TSCA Section 8(e) Compliance Audit Program (CAP) should already contain information supporting confidentiality claims, because substantiation of CBI claims is required at the same time the 8(e) CAP is submitted to EPA. (If not done so already, please ensure that this information is provided to the Agency). When substantiating any/all claims, answer the questions detailed in the following attachment.

For NON-CAP submissions, any confidentiality claims should be supported by submission of information as described in the attachment(s).

12199 A



CECATS DATA:

Submission # BEHQ 0893-12/99 SEQ. A

TYPE: INT. SUPP FLWP

SUBMITTER NAME: Dow Chemical
Company

SUB. DATE: 08/31/93 OTS DATE: 09/03/93 CSRAD DATE: 09/22/93

CHEMICAL NAME:

Diethylene Triamine

INFORMATION REQUESTED: FLWP DATE:

0501 NO INFO REQUESTED
0502 INFO REQUESTED (TECH)
0503 INFO REQUESTED (VOL ACTIONS)
0504 INFO REQUESTED (REPORTING RATIONALE)
DISPOSITION:
0670 REFER TO CHEMICAL SCREENING
0678 CAP NOTICE

VOLUNTARY ACTIONS:

0401 NO ACTION REPORTED
0402 STUDIES PLANNED/UNDERWAY
0403 NOTIFICATION OF WORK/RA/OTHERS
0404 LABEL/MSDS CHANGES
0405 PROCESS/HANDLING CHANGES
0406 APP/USE DISCONTINUED
0407 PRODUCTION DISCONTINUED
0408 CONFIDENTIAL

CASE#

001-111-40-0

INFORMATION TYPE:

P F C

0201	ONCO (HUMAN)	01 02 04
0202	ONCO (ANIMAL)	01 02 04
0203	CELL TRANS (IN VITRO)	01 02 04
0204	MUTA (IN VITRO)	01 02 04
0205	MUTA (IN VIVO)	01 02 04
0206	REPRO/TERATO (HUMAN)	01 02 04
0207	REPRO/TERATO (ANIMAL)	01 02 04
0208	NEURO (HUMAN)	01 02 04
0209	NEURO (ANIMAL)	01 02 04
0210	ACUTE TOX (HUMAN)	01 02 04
0211	CHR. TOX (HUMAN)	01 02 04
0212	ACUTE TOX (ANIMAL)	01 02 04
0213	SUB ACUTE TOX (ANIMAL)	01 02 04
0214	SUB CHRONIC TOX (ANIMAL)	01 02 04
0215	CHRONIC TOX (ANIMAL)	01 02 04

INFORMATION TYPE:

0216	EPICLIN	01 02 04
0217	HUMAN EXPOS (PROD CONTAM)	01 02 04
0218	HUMAN EXPOS (ACCIDENTAL)	01 02 04
0219	HUMAN EXPOS (MONITORING)	01 02 04
0220	BIOAQUA TOX	01 02 04
0221	ENV. OCCURRENCE/FATE	01 02 04
0222	EMER INCI OF ENV CONTAM	01 02 04
0223	RESPONSE REQUEST DELAY	01 02 04
0224	PROD/COMP/CHEM ID	01 02 04
0225	REPORTING RATIONALE	01 02 04
0226	CONFIDENTIAL	01 02 04
0227	ALLERG (HUMAN)	01 02 04
0228	ALLERG (ANIMAL)	01 02 04
0229	METAB/PHARMACO (ANIMAL)	01 02 04
0230	METAB/PHARMACO (HUMAN)	01 02 04

P F C

INFORMATION TYPE:

0241	IMMUNO (ANIMAL)	01 02 04
0242	IMMUNO (HUMAN)	01 02 04
0243	CHEMPHYS PROP	01 02 04
0244	CLASTO (IN VITRO)	01 02 04
0245	CLASTO (ANIMAL)	01 02 04
0246	CLASTO (HUMAN)	01 02 04
0247	DNA DAM/REPAIR	01 02 04
0248	PROD/USE/PROC	01 02 04
0251	MSDS	01 02 04
0299	OTHER	01 02 04

P F C

TRIAGE DATA: NON-CR INVENTORY

YES (CONTINUE)

NO (DROP)

NO CAP

ONGOING REVIEW

YES (DROP/REFE.)

NO (CONTINUE)

REFER.

SPECIES

RAT

TOXICOLOGICAL CONCERN

LOW

MED M - study not complete
effects at 300 mg/kg &
perhaps at 100 mg/kg

HIGH

USE:

PRODUCTION:

COMMENTS: Non-CAP